

(19) World Intellectual Property
Organization
International Bureau



(43) International Publication Date
1 July 2004 (01.07.2004)

PCT

(10) International Publication Number
WO 2004/054980 A1

(51) International Patent Classification⁷: **C07D 217/26**

(74) Agents: **MAJUMDAR, S. et al.**; S. Majumdar & Co., 5, Harish Mukherjee Road, Calcutta 700 025 (IN).

(21) International Application Number:
PCT/IN2002/000235

(22) International Filing Date:
16 December 2002 (16.12.2002)

(25) Filing Language: English

(26) Publication Language: English

(71) Applicant (for all designated States except US): **LUPIN LIMITED** [IN/IN]; 159 C.S.T. Road, Kalina, Santacruz (East), Mumbai 400 098, Maharashtra (IN).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **SINGH, Girij, Pal** [IN/IN]; Lupin Limited (Research Park), 46A/47A, Nande Village, Taluka Mulshi, Pune 411 042, Maharashtra (IN). **RAWAT, Govind, Singh** [IN/IN]; Lupin Limited (Research Park), 46A/47A, Nande Village, Taluka Mulshi, Pune 411 042, Maharashtra (IN). **DHAKE, Vilas, Nathu** [IN/IN]; Lupin Limited (Research Park), 46A/47A, Nande Village, Taluka Mulshi, Pune 411 042, Maharashtra (IN). **NEHATE, Sagar, Purshottam** [IN/IN]; Lupin Limited (Research Park), 46A/47A, Nande Village, Taluka Mulshi, Pune 411 042, Maharashtra (IN).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.

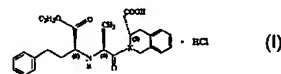
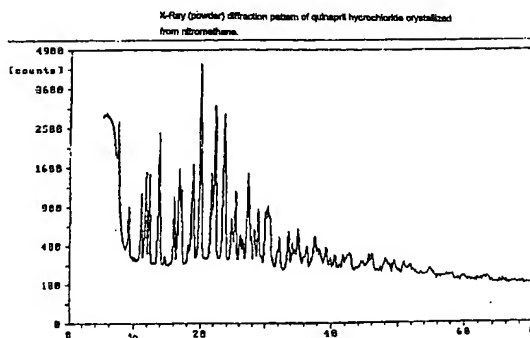
(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: **CRYSTALLINE FORM OF QUINAPRIL HYDROCHLORIDE AND PROCESS FOR PREPARING THE SAME**



(57) Abstract: A novel crystalline form of quinapril hydrochloride of formula (I). An amorphous form of quinapril hydrochloride substantially free of impurities, specially diketopiperazine compound, and conforming to pharmacopoeial specifications formed from the said novel crystalline form of quinapril hydrochloride of formula (I). The crystalline quinapril hydrochloride is in the form nitroalkane solvate in which the nitroalkane is nitromethane, nitroethane and nitropropane. Each such nitroalkane solvate having particular characteristic X- ray diffraction patterns. A process for preparation of amorphous from of quinapril hydrochloride, substantially free of impurities, specially diketopiperazine compound, and conforming to pharmacopoeial specifications, using the novel crystalline quinapril hydrochloride as an intermediate. The process involves obtaining free base compound of formula (V) by adjusting the pH of a solution of the benzyl ester maleate salt of quinapril of formula (V) between 7.5-8.5 in a mixture of water and an organic solvent; catalytic hydrogenation of this compound (V) in an alcoholic solvent in the presence of concentrated hydrochloric acid or hydrogen chloride dissolved in an alcoholic solvent and in the presence of catalytic amounts of Pd/C to obtain a residue containing formula (I); crystallization of the said residue by evaporating the alcoholic solvent from a nitroalkane solvent to give crystalline quinapril hydrochloride, associated with a solvate of the nitroalkane solvent, and drying the crystalline quinapril hydrochloride nitroalkane solvate at a temperature between 40°C and 45°C under vacuum to give amorphous quinapril hydrochloride of formula (I).

EXPRESS MAIL NO.: ER188423891US
DOCKET NO.: PC25242A

WO 2004/054980 A1

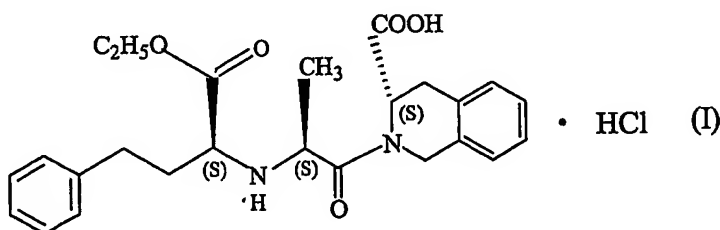
CRYSTALLINE FORM OF QUINAPRIL HYDROCHLORIDE AND PROCESS FOR PREPARING THE SAME

FIELD OF THE INVENTION

The present invention relates to a novel crystalline form of quinapril hydrochloride of formula (I) and a process for its preparation thereof. The present invention further relates to the preparation of amorphous form of quinapril hydrochloride, of high purity and conforming to pharmacopoeial specifications, by the use of the novel crystalline quinapril hydrochloride of formula (I) as an intermediate.

BACKGROUND OF THE INVENTION

The chemical species, (3S)-2-[(2S)-2-[(1S)-1-(Ethoxycarbonyl)-3-phenylpropyl]amino]-1-oxopropyl]-1,2,3,4-tetrahydro-3-isoquinolinecarboxylic acid is known generically as quinapril. Its pharmaceutically acceptable salts, specially the hydrochloride salt is represented by formula (I).

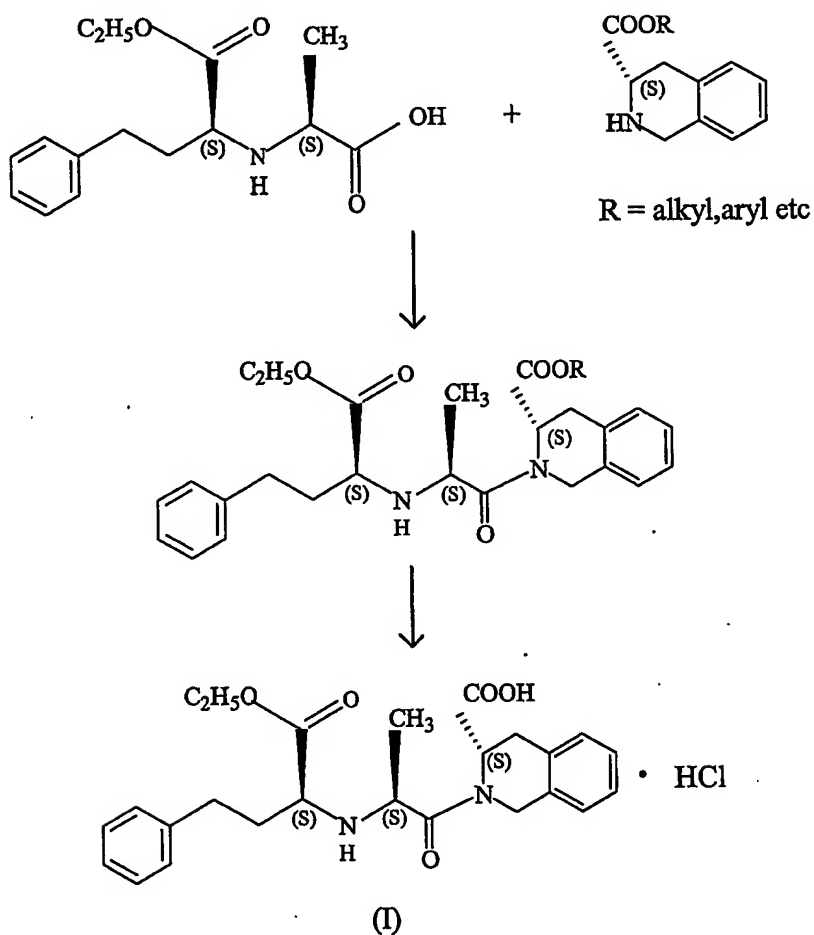


Quinapril hydrochloride (I) and its pharmaceutically acceptable salts are active as angiotensin converting enzyme (ACE) inhibitors and thus are commercially valuable antihypertensive agents.

US Patent No. 4,344,949 (Hoeftle et. al.) *inter alia* describes a process for the preparation of quinapril hydrochloride comprising reaction of ethyl ester of (S,S)- α -[(1-carboxyethyl)amino] phenylbutanoic acid with the benzyl or t-butyl ester of (S)-1,2,3,4-tetrahydro-3-isoquinoline carboxylic acid in the presence of 1-hydroxy benzotriazole, employing standard peptide bond forming methods.

The benzyl or t-butyl ester group of quinapril thus obtained is removed by catalytic hydrogenation or by treatment with trifluoroacetic acid. Quinapril hydrochloride is isolated either by solvent precipitation with diethyl ether or by lyophilisation of the aqueous solution.

5 The above method is summarised in scheme (I).

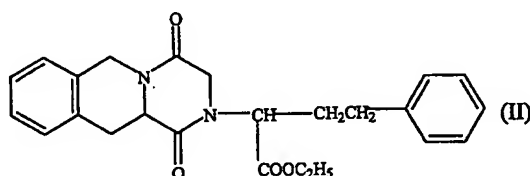


Scheme-I : Synthesis of quinapril hydrochloride as disclosed in US Patent No. 4,344,949

10 Other methods for preparation of quinapril hydrochloride are disclosed by Barton et. al. in GB Patent No. 2,095,252, by Patchett et. al. in EP Patent No. 0,065,301 etc.

However, all the reported methods for synthesis of quinapril suffer from a serious drawback in that the product obtained by all the methods is invariably contaminated with varying amounts of an impurity identified as the diketopiperazine derivative of formula (II), leading, most often to product not conforming to pharmacopoeial specifications.

- 5 The diketopiperazine impurity is formed either during removal of the carboxylic acid protective group or it could be formed during drying of quinapril hydrochloride.



The above impurity once formed is difficult to remove by conventional separation techniques, including fractional crystallization.

- 10 Regulatory authorities all over the world are becoming very stringent about the purity of an approved drug or a drug candidate awaiting approval. Especially there is a growing concern about the nature and level of impurities present in such molecules.

- 15 Pharmaceutical manufacturers throughout the world, constantly strive to produce drug molecules that go beyond pharmacopoeial specifications i.e. compounds having extra pharmacopoeial specifications.

US Patent No. 4,761,479 (Goel et. al.) discloses a method wherein quinapril hydrochloride obtained from the reaction mixture is crystallized from a solvent selected from acetonitrile or acetone to give a crystalline form of quinapril hydrochloride possessing a characteristic X-ray (powder) diffraction pattern and high bulk density.

- 20 The abovementioned patent also mentions that the crystalline material thus obtained contains equimolar amounts of acetonitrile/acetone as part of the crystal lattice. The patent further mentions that the solvent(s) present in the crystal structure can be removed under vacuum at a temperature of about 50⁰ C. However during this unit operation the crystallinity of the substance is lost due to desolvation during drying, and an amorphous

material is obtained which is claimed to be free of impurities, specially the diketopiperazine compound of formula (II).

Even though, the US Patent No. 4,761,479 discloses the use of acetonitrile and acetone for crystallization, the former is preferred since, unlike acetonitrile, acetone cannot be removed from the crystal lattice, even on prolonged drying.

Canadian Application No. CA 12,293,705 A1 (Llagostera et. al) describes a process for purification of impure quinapril hydrochloride through a two-step process, comprising first crystallisation of the impure material from toluene followed by another crystallization using a class III solvent as categorized by International Conference of Harmonisation (ICH). The patent also mentions that initially a solvate of quinapril hydrochloride with toluene is formed, which is subsequently substituted by a solvate of the class III solvent used for the second crystallisation. This patent like the aforesaid US Patent No. 4,761,479 also claims that amorphous quinapril hydrochloride is obtained on desolvation of the solvate of the material with the class III solvent through drying. Among the class III solvents disclosed, ethyl formate and methyl acetate are the preferred ones.

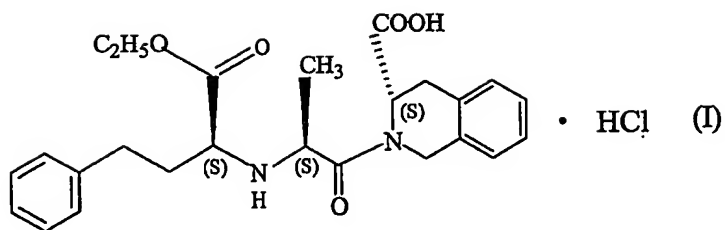
However, the abovementioned two methods for purification of impure quinapril hydrochloride utilise solvents which have low flash points eg. the flash point of acetonitrile is 2⁰C, that of methyl acetate is -16⁰ C, whereas that of ethyl formate is -20⁰ C, rendering their use on commercial scale a hazardous proposition. Moreover, the method disclosed in the later patent i. e. Canadian Application No. 2,293,705 A1 involves a two-step purification, which makes the entire operation not only time consuming but costly.

The need, therefore, exists for a method for obtaining quinapril hydrochloride of high purity which addresses the shortcomings associated with the prior art methods.

25

SUMMARY OF THE INVENTION

According to the principal aspect of present invention there is provided a pure crystalline quinapril hydrochloride of formula I associated with the solvate of the nitroalkane.



According to further aspect of the present invention there is provided a novel crystalline nitromethane solvate of quinapril hydrochloride of formula I exhibiting essentially the following X-ray (powder) diffraction properties:

5

Spacing 'd'	Relative intensity
16.2471	63.5
13.8426	55.5
11.9609	57.3
9.6467	17.2
7.9468	22.4
7.5064	31.1
7.1699	30.2
6.4095	52.4
6.0561	3.1
5.5041	20.4
5.2808	33.1
5.1767	22.8
4.8704	5.7
4.6830	34.4
4.4404	100
4.0977	30.9
3.9931	69.5
3.7747	62.8
3.5972	12.9
3.5058	22.8
3.4153	8.3

3.3558	7.0
3.2676	30.7
3.2054	7.6
3.1510	9.5
3.0854	16.5
2.9772	14.5
2.9403	17.5
2.9122	12.9
2.7798	7.6
2.6670	9.0
2.6216	6.0
2.5613	10.1
2.4650	5.2
2.3933	7.9
2.2963	4.9
2.2620	2.7
2.2290	3.0
2.1672	3.5
2.1125	3.4
2.0361	2.0
1.9911	3.3
1.9714	3.4
1.8935	2.5
1.8420	2.3
1.7917	2.0
1.7630	1.3
1.6723	1.1
1.5928	0.4
1.4683	0.5

According to another aspect of the present invention there is provided a novel crystalline nitroethane solvate of quinapril hydrochloride of formula I exhibiting essentially the following X-ray (powder) diffraction properties:

5

Spacing 'd'	Relative intensity
17.4844	87.0
16.0841	67.2
12.0996	53.3
10.0860	18.8
8.1700	29.6
7.7522	41.2
7.3814	35.4
6.5032	39.9
6.0580	2.9
5.5780	35.8
5.3973	32.0
5.2776	23.7
4.9335	19.6
4.8335	32.9
4.7627	40.5
4.5635	88.9
4.5095	69.6
4.4227	36.5
4.2021	36.0
4.0818	100.0
3.8719	30.4
3.7802	48.7
3.6435	49.5
3.4889	10.4
3.3520	20.8

3.2948	36.3
3.1526	15.3
3.1132	19.1
3.0304	49.5
2.9280	19.2
2.9080	14.1
2.7933	7.9
2.6986	10.8
2.6399	9.4
2.5715	16.7
2.5194	9.6
2.4535	4.6
2.4140	12.7
2.3567	8.5
2.3093	5.5
2.2801	6.1
2.1687	6.7
2.1303	4.5
2.0332	5.0
2.0031	6.6
1.9801	3.7
1.9395	2.7
1.8918	4.2
1.8632	3.5
1.8354	3.0
1.8110	2.4
1.7812	2.1
1.7024	1.1
1.5414	0.6
1.3867	0.4

According to another aspect of the present invention there is provided a novel crystalline nitropropane solvate of quinapril hydrochloride of formula I exhibiting essentially the following X-ray (powder) diffraction properties

5

Spacing 'd'	Relative intensities
17.4327	100.0
15.4378	65.5
12.1079	49.5
10.4492	18.2
8.3626	21.3
7.9183	38.9
7.5253	27.5
6.6268	19.6
6.4583	17.2
5.6238	43.2
5.2713	27.8
5.1011	18.3
4.9816	30.7
4.8049	38.7
4.6878	52.0
4.5427	61.4
4.4480	35.3
4.3008	25.3
4.1690	68.6
4.0279	14.7
3.9483	23.9
3.7993	57.5
3.5871	15.5
3.4582	14.8
3.2960	32.0

3.1558	20.6
3.1053	26.1
3.0548	17.9
2.9559	9.2
2.9066	9.9
2.8044	10.6
2.7274	6.5
2.6357	9.4
2.5910	11.1
2.4880	5.3
2.4365	6.4
2.4053	6.0
2.3446	8.4
2.3084	5.2
2.2682	4.1
2.2075	3.0
2.1667	4.1
2.0297	13.7
2.0023	3.5
1.8765	2.2
1.8019	2.6
1.4337	2.2
1.4005	0.4

According to a further aspect of the present invention there is provided an amorphous form of quinapril hydrochloride of formula I, in high purity, free of impurities and conforming to pharmacopoeial specifications obtained from the novel crystalline form of quinapril hydrochloride associated with the solvate of the nitroalkane.

5

Thus according to a still further aspect of the present invention there is provided a simple, industrial method for preparation of quinapril hydrochloride of formula I of high purity.

According to another aspect of the present invention there is provided a simple and industrial method for purification of quinapril hydrochloride of formula I comprising
5 crystallisation of impure quinapril hydrochloride from a nitroalkane solvent to give a novel crystalline form of quinapril hydrochloride associated with a solvate of the solvent utilised for crystallisation.

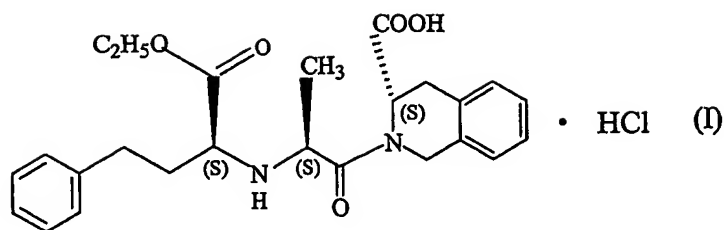
According to further aspect of the present invention there is provided a method for obtaining an amorphous form of quinapril hydrochloride, of high purity, free of impurities
10 and conforming to pharmacopoeial specifications from the novel crystalline form of quinapril hydrochloride associated with the solvate of the nitroalkane solvent.

According to further aspect of the present invention there is provided a method for preparation of amorphous form of quinapril hydrochloride, in high purity, free of impurities and conforming to pharmacopoeial specifications comprising removal of the
15 solvent from the novel crystalline form of quinapril hydrochloride associated with the solvate of the nitroalkane solvent by subjecting the said crystalline form to drying under vacuum.

According to another aspect of the present invention there is provided quinapril hydrochloride of formula I obtained by a simple and industrial method for purification of
20 quinapril hydrochloride comprising crystallisation of impure quinapril hydrochloride from a nitroalkane solvent to give a novel crystalline form of quinapril hydrochloride associated with a solvate of the solvent utilised for crystallisation.

DETAILED DESCRIPTION OF THE INVENTION

25 According to the present invention, quinapril hydrochloride of formula I



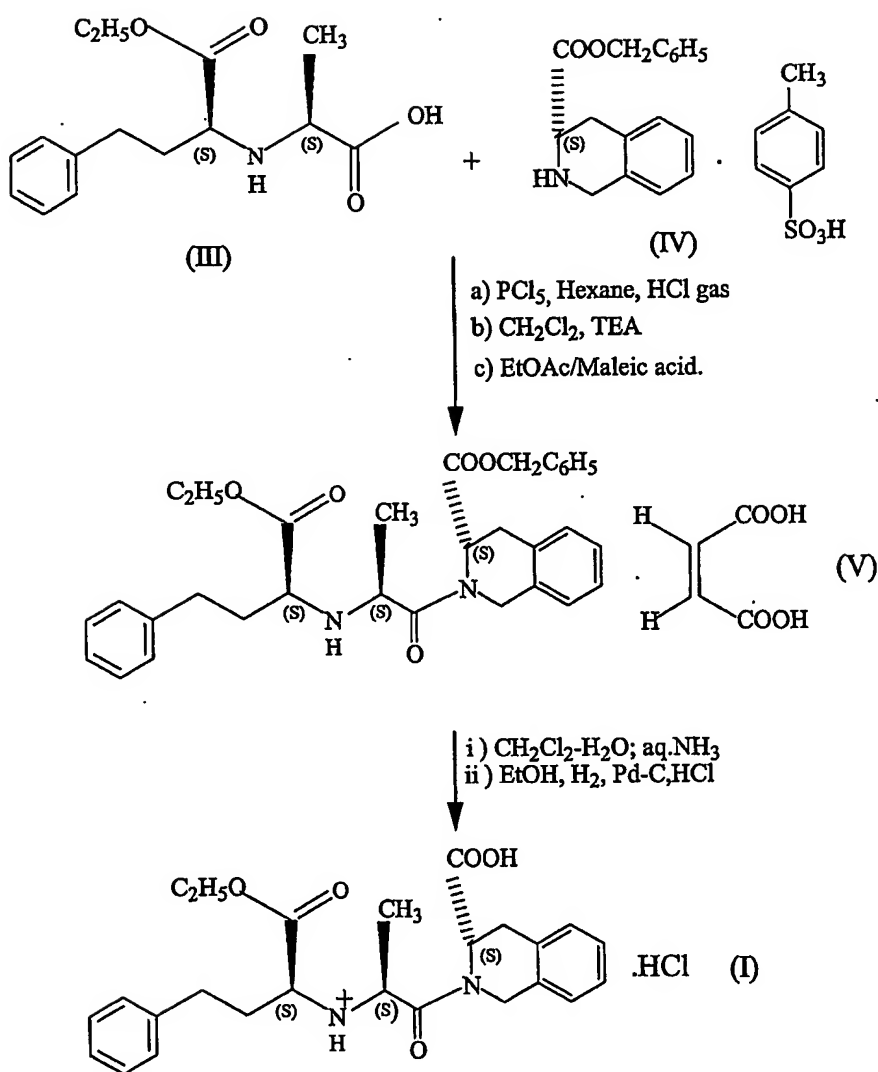
can be synthesized by two methods, as summarized in Method-I and Method-II.

5 Method-I:

The first step of Method-I consists of condensation of N-[1(S)-carboethoxy-3-phenylpropyl]-(S)-alanine (III) with the para-toluene sulfonate salt of the benzyl ester of (S)-1,2,3,4-Tetrahydroisoquinoline-3-carboxylic acid (IV) to give the benzyl ester of (S, S, S)-2-{2-[(1-ethoxycarbonyl)-3-phenylpropyl]amino}-1-oxopropyl]-1,2,3,4-tetrahydro-
 10 3-isoquinoline carboxylic acid or quinapril benzyl ester which is isolated as the maleic acid salt (V).

Typically, N-[1(S)-carboethoxy-3-phenylpropyl]-(S)-alanine (III) is converted into its acid chloride by reacting with either PCl_5 or PCl_3 or SOCl_2 , but preferably with PCl_5 in a
 15 non-polar or polar solvent, such as those selected from chlorinated hydrocarbons like dichloromethane or 1,2-dichloroethane; aromatic hydrocarbons like toluene, xylene or aliphatic hydrocarbons like hexane, heptane etc. A non-polar solvent is preferred as the acid chloride precipitates out from such solvents and can be easily isolated by filtration.

13



Method-I: Preparation of quinapril hydrochloride

The acid chloride is prepared by treating the acid (III) with PCl_5 , using hexane as a solvent, the molar ratio of PCl_5 used being in the range of 0.9 moles to 1.5 moles, but preferably in the range of 1.0 mole to 1.2 moles.

The reaction temperature employed can be between -5°C to $+15^\circ\text{C}$ and for a period of time between 2-5 hrs, but preferably the temperature is in the range of $0\text{-}10^\circ\text{C}$ and reaction time between 3-4 hours.

The acid chloride thus obtained is then condensed with the free base of the benzyl ester of (S)-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid (IV). The free base is obtained by treatment of the organic or inorganic acid salt of (S)-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid (IV) with an organic or inorganic base like trialkyl amine, alkali hydroxide etc.

Preparation of compound (V) is then effected in chlorinated solvents like dichloromethane, 1,2-dichloroethane, chloroform etc comprising by addition of the acid chloride of compound (III) to the solution of compound (IV) in the presence of an organic base, such as selected from triethylamine, diethylamine, tertiary butylamine, and dicyclohexylamine at temperatures ranging from -30°C to 0°C , but preferably at -15 to 20°C and subsequently raising the temperature in the range of 10 to 40°C , but preferably 20 to 30°C . The compound (V) is isolated by washing the organic layer with water and formation of its salt with an organic or an inorganic acid.

In a specific embodiment, a solution of the acid (III) in hexane is reacted with stoichiometric amounts of phosphorous pentachloride at a temperature between 0 to 10°C , with agitation for 3 hrs. The acid chloride thus formed is isolated by filtration.

A solution of the acid chloride thus obtained is dissolved in dichloromethane and added to the free base of compound (IV). The free base of (IV) is obtained by treating the corresponding p-toluenesulfonate salt with an equimolar amount of organic base, like triethyl amine.

The condensation between the free base of compound (IV) and the acid chloride of compound (III) is carried out in a chlorinated solvent eg. dichloromethane. The acid chloride is added to the free base at -15°C in the presence of an equimolar amount of an organic base eg. triethyl amine. The reaction is then carried forward to completion at room temperature by agitating for a period ranging from 30 to 90 minutes. The reaction mixture is worked up by successively washing the reaction mixture with hydrochloric acid, followed by water and finally with aqueous sodium bicarbonate solution.

Quinapril benzyl ester thus obtained is isolated as its maleate salt by treating the quinapril benzyl ester with maleic acid at room temperature in a alkyl acetate solvent, preferably ethyl acetate.

The second step comprises hydrogenolysis of quinapril benzyl ester maleate salt (V) thus
5 obtained in the presence of Pd/C catalyst and concentrated hydrochloric acid.

The hydrogenolysis of the benzyl ester (V) can be carried out in an alcoholic solvent, such as methanol, ethanol or isopropanol in presence of concentrated hydrochloric acid or a solution of hydrogen chloride in the same alcohol, with hydrogen gas at a pressure
10 ranging between 10-70 psi and at a temperature between 10-40⁰ C.

In a specific embodiment, the hydrogenolysis reaction is carried out on the free base of quinapril benzyl ester (V), which in turn is obtained from the maleate salt (V) by treatment with aqueous ammonia at slightly alkaline pH of 7.5-8.5. Hydrogenolysis is
15 carried out using ethanol as a solvent and in presence of concentrated hydrochloric acid at a pressure ranging between 40-60 psi and room temperature.

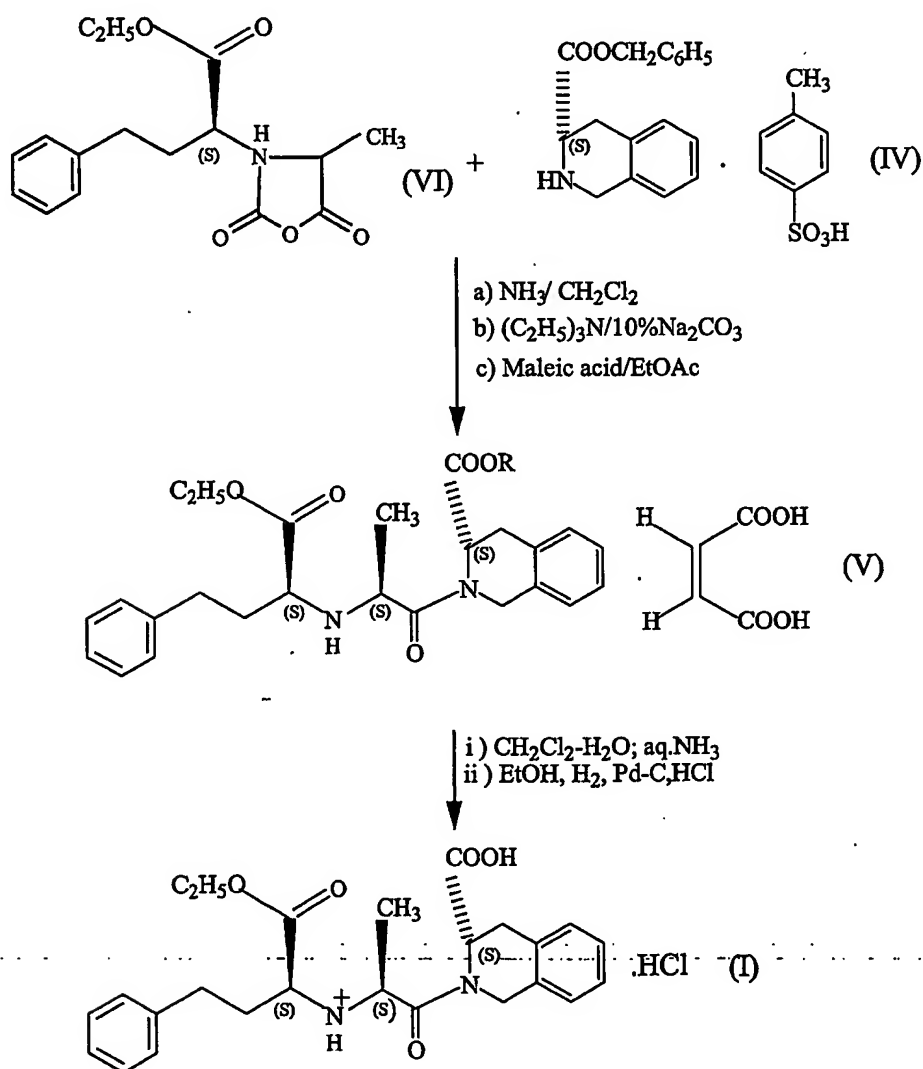
The molar ratio between the benzyl ester of quinapril (V) and hydrochloric acid can be equal or greater than the stoichiometric ratio (larger amount of hydrochloric acid in the
20 reaction mixture leads to increase in the diketopiperazine impurity (II), arising due to cyclisation of quinapril hydrochloride under acidic conditions).

The catalyst is removed by filtration and the filtrate evaporated at low temperature, preferably below 30⁰ C as higher temperature could lead to formation of more amounts of
25 the diketopiperazine impurity (II).

Method-II:

The second method comprises of reacting N [1(S)-carboethoxy-3-phenylpropyl]-(S)-alanyl carboxyanhydride (VI) with the free base of the benzyl ester of (S)-1,2,3,4-
30 tetrahydroisoquinoline-3-carboxylic acid (IV).

The free base of (IV) is obtained by treating the organic or inorganic acid salt of (S)-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid (IV) with an organic or inorganic base like trialkyl amine, alkali hydroxide aqueous ammonia etc in a mixture of water and chlorinated solvents like 1,2-dichloroethane, dichloromethane chloroform etc. The pH of the mixture is adjusted between 8.0 -9.0 at a temperature between -15 to +15°C, but preferably between 0-10°C. The reaction mixture is then stirred at room temperature, for 15 to 45 minutes, but preferably between 25-35minutes. The organic layer containing the free base is separated.



Method-II: Preparation of quinapril hydrochloride

The compound, N [1(S)-carboethoxy-3-phenylpropyl]-(S)-alanyl carboxyanhydride (VI) is dissolved in chlorinated solvents like dichloromethane, 1,2-dichloroethane or chloroform, but preferably in dichloromethane at room temperature. This mixture is then added to the free base of (IV) dissolved in dichloromethane, at a temperature between 20 to 45°C, but preferably at 25 to 35°C. The reaction mixture is agitated for a period between 1 to 5 hours, but preferably between 2 to 4 hours for the reaction to go to completion. A mixture of an aqueous solution of an inorganic base like sodium bicarbonate, sodium carbonate and an organic base like triethyl amine, dicyclohexyl amine etc is added to neutralize the reaction mixture. The organic layer is separated and concentrated and the residue is diluted with an alkyl acetate preferably ethyl acetate. The resultant mixture is stirred at a temperature between 30 to 50°C, but preferably between 35 to 45°C. The product is isolated as a maleate salt as described in method -I.

In a specific embodiment, (S)-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid (IV), is dissolved in a mixture of water and dichloromethane at 5 to 10°C, and treated with aqueous ammonia solution, at the same temperature. The organic layer is separated and treated with N[1(S)-carboethoxy-3-phenylpropyl]-(S)-alanyl carboxyanhydride (VI), dissolved in dichloromethane at 25-35°C, the resultant mixture is stirred for 2-4 hours, at the same temperature till the reaction goes to completion. The reaction mixture is then neutralized with a mixture of aqueous sodium carbonate and triethyl amine and stirred for 2-4 hours. The organic layer is separated and concentrated, the residue is diluted with ethyl acetate. The mixture is then treated with a mixture of maleic acid dissolved in ethyl acetate to obtain benzyl quinapril ester as the maleate salt as described in method-I.

The benzyl ester quinapril maleate salt is then converted by catalytic hydrogenolysis to quinapril hydrochloride, as in method -I.

The residue of quinapril hydrochloride (I) thus obtained is made anhydrous by addition of dry dichloromethane. The resulting mixture is distilled to remove water as an azeotrope. This procedure is repeated until the moisture content of the mixture is less than 1.0%.

The amount of the diketopiperazine impurity (II) present in the quinapril hydrochloride thus obtained varies between 2.0-5.0% by weight of the quinapril hydrochloride.

Purification of the impure quinapril hydrochloride thus obtained is effected by crystallisation from a nitroalkane solvent from which a crystalline form of quinapril hydrochloride falls out. The nitroalkane solvent is selected from nitromethane, nitroethane, and nitropropane. Among these, nitromethane is preferred since it is less
5 costly and readily available.

The nitroalkane solvents like acetonitrile belong to Class II solvents as categorised by the International Conference on Harmonisation (ICH). However, unlike acetonitrile these solvents have higher flash point eg. that of nitromethane being 44°C as compared to 2°C for acetonitrile, -16°C for methyl acetate and -20°C for ethyl formate, rendering them
10 easier and safer to handle during commercial manufacture.

The crystallisation is carried out by addition of the nitroalkane solvent to the anhydrous residue of impure quinapril hydrochloride, obtained as per the method described hereinearlier at room temperature. The mixture is stirred for 10-15 minutes to get a clear solution. The solution is seeded with pure quinapril hydrochloride and then cooled to a
15 temperature ranging between -15 to $+15^{\circ}\text{C}$, preferably in the range of 0 to $+10^{\circ}\text{C}$ and agitated at the same temperature for a period of 1-3 hours, preferably 2.0 hrs, to allow the nitroalkane solvate of quinapril hydrochloride to crystallize out. The crystalline solid is filtered out at room temperature.

In a specific embodiment, the impure quinapril hydrochloride is dissolved in
20 nitromethane at room temperature. The amount of nitromethane used can be between 5-10 times by volume, of the weight of the impure material. The solution is seeded with pure quinapril hydrochloride at same temperature and cooled to $5-10^{\circ}\text{C}$. The crystalline solvate is allowed to crystallize out at a temperature between $5-10^{\circ}\text{C}$ and the crystalline material filtered off at room temperature.

25 The quinapril hydrochloride to nitroalkane solvate ratio was determined on the basis of ^1H NMR & HPLC and was found to be having a equimolar ratio.

These novel crystalline solvates quinapril hydrochloride with the nitroalkane solvents possess distinct X-ray (powder) diffraction patterns and these are summarized in Table-I.

Table-I: X-ray (powder) diffraction pattern of the nitroalkane solvates of quinapril hydrochloride

Nitromethane		Nitroethane		Nitropropane	
Spacing 'd'	Relative intensity	Spacing 'd'	Relative intensity	Spacing 'd'	Relative intensity
16.247	63.5	17.4844	87.0	17.4327	100.0
13.8426	55.5	16.0841	67.2	15.4378	65.5
11.9609	57.3	12.0996	53.3	12.1079	49.5
9.6467	17.2	10.0860	18.8	10.4492	18.2
7.9468	22.4	8.1700	29.6	8.3626	21.3
7.5064	31.1	7.7522	41.2	7.9183	38.9
7.1699	30.2	7.3814	35.4	7.5253	27.5
6.4095	52.4	6.5032	39.9	6.6268	19.6
6.0561	3.1	6.0580	2.9	6.4583	17.2
5.5041	20.4	5.5780	35.8	5.6238	43.2
5.2808	33.1	5.3973	32.0	5.2713	27.8
5.1761	22.8	5.2776	23.7	5.1011	18.3
4.8704	5.7	4.9335	19.6	4.9816	30.7
4.6830	34.4	4.8335	32.9	4.8049	38.7
4.4404	100	4.7627	40.5	4.6878	52.0
4.0977	30.9	4.5635	88.9	4.5427	61.4
3.9931	69.5	4.5095	69.6	4.4480	35.3
3.7747	62.8	4.4227	36.5	4.3008	25.3
3.5972	12.9	4.2021	36.0	4.1690	68.6
3.5058	22.8	4.0818	100.0	4.0279	14.7

Table-I (Continued)

Nitromethane		Nitroethane		Nitropropane	
Spacing 'd'	Relative intensity	Spacing 'd'	Relative intensity	Spacing 'd'	Relative intensity
3.4153	8.3	3.8719	30.4	3.9483	23.9
3.3558	7.0	3.7802	48.7	3.7993	57.5
3.2676	30.7	3.6435	49.5	3.5871	15.5
3.2054	7.6	3.4889	10.4	3.4582	14.8
3.1510	9.5	3.3520	20.8	3.2960	32.0
3.0854	16.5	3.2948	36.3	3.1558	20.6
2.9772	14.5	3.1526	15.3	3.1053	26.1
2.9403	17.5	3.1132	19.1	3.0548	17.9
2.9122	12.9	3.0204	49.5	2.9559	9.2
2.7798	7.6	2.9280	19.2	2.9066	9.9
2.6670	9.0	2.9080	14.1	2.8044	10.6
2.6216	6.0	2.7933	7.9	2.7274	6.5
2.5613	10.1	2.6986	10.8	2.6357	9.4
2.4650	5.2	2.6399	9.4	2.5910	11.1
2.3933	7.9	2.5715	16.7	2.4880	5.3
2.2963	4.9	2.5194	9.6	2.4365	6.4
2.2620	2.7	2.4535	4.6	2.4053	6.0
2.2290	3.0	2.4140	12.7	2.3446	8.4

Table-I (Continued)

Nitromethane		Nitroethane		Nitropropane	
Spacing 'd'	Relative intensity	Spacing 'd'	Relative intensity	Spacing 'd'	Relative intensity
2.1672	3.5	2.3567	8.5	2.3084	5.2
2.1125	3.4	2.3093	5.5	2.2682	4.1
2.0361	2.0	2.2801	6.1	2.2075	3.0
1.9911	3.3	2.1687	6.7	2.1667	4.1
1.9714	3.4	2.1303	4.5	2.0297	13.7
1.8935	2.5	2.0332	5.0	2.0023	3.5
1.8420	2.3	2.0031	6.6	1.8765	2.2
1.7917	2.0	1.9801	3.7	1.8019	2.6
1.7630	1.3	1.9395	2.7	1.4337	2.2
1.6723	1.1	1.8918	4.2	1.4005	0.4
1.5928	0.4	1.8632	3.5	-----	-----
1.4683	0.5	1.8354	3.0		
-----	-----	1.8110	2.4		
		1.7812	2.1		
		1.7024	1.1		
		1.5414	0.6		
		1.3867	0.4		
		-----	-----		

All the X-ray diffraction analysis were carried out by the powder diffraction method ($\lambda=1.54060$), the preparations of the sample were performed on a dry standard.

Diffractometer type: PW1710 BASED

Material of the anode: Copper.

Wavelength alpha1 [λ] = 1.54060

Wavelength alpha 2 [λ] :1.54439

Initial angle [$2\theta^\circ$]: 5.010

Final angle [$2\theta^\circ$]; 69.810

5 Intensity ratio (alpha2/alpha1): 0.500

Maximum intensity: 3893.760

The X-ray (powder) diffraction pattern of the crystalline nitromethane, nitroethane and nitropropane solvates are reproduced in Fig Ia, Ib and Ic respectively.

10 The I.R. spectra of the crystalline nitromethane, nitroethane and nitropropane solvates are reproduced in Fig IIa, IIb and IIc respectively.

Subsequent drying of the crystalline solvate at $40^\circ\text{C}/5\text{-}10\text{mmHg}/60\text{-}70$ hours, desolvates the crystalline compound to give amorphous quinapril hydrochloride of high purity which conforms to pharmacopoeial specifications. The crystalline solvate is typically dried at 40°C under vacuum at $5\text{-}10\text{mmHg}$ for $60\text{-}70$ hours.

15

The quinapril thus obtained after drying is substantially free from impurities, specially the diketopiperazine compound of formula (II) and easily conforms to pharmacopoeial specifications and highly suitable for human use.

20 The invention can be further illustrated by the following examples, which, however, should not be construed as limiting the scope of the invention.

Example-1

Preparation of the benzyl ester of (S, S, S)-2-[2-[(1-ethoxycarbonyl)-3-phenylpropyl)amino]-1-oxopropyl]-1,2,3,4-tetrahydro-3-isoquinoline carboxylic acid; maleic acid salt (Quinapril benzyl ester maleic acid salt). (Method-I)

25

25gms (0.089 moles) of N[1(S)-carboethoxy-3-phenylpropyl]-(S)-alanine (III) was taken in hexane, cooled to -5°C . To this was added 20.5 gms of phosphorous pentachloride in

lots, followed by purging of hydrogen chloride gas. The reaction mixture was stirred at 5-10⁰ C for 3hrs. The acid chloride, which precipitated out, was filtered and washed with n-hexane.

5 35.4gms (0.086 moles) of the para-toluene sulfonate salt of the benzyl ester of (S)-1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid (IV) was dissolved in 150ml dichloromethane. 29ms (0.287 moles) of triethyl amine was added at 0-5⁰ C and the reaction mixture stirred at 0-5⁰ C for 10minutes. A solution of the acid chloride of compound (III) prepared in the earlier step in 150ml of dichloromethane was added at -15
10 to -20⁰ C to the reaction mixture. The reaction mixture was then stirred at 20-25⁰ C for 1hour. The organic phase was separated and washed with 200ml of hydrochloric acid followed by 200ml of water. The organic phase was then concentrated under vacuum. The residue of the quinapril benzyl ester was dissolved in 130ml ethyl acetate and washed with 200ml of 10% aqueous sodium bicarbonate solution followed by water wash
15 (200ml).

A solution of 9.4 gms of maleic acid dissolved in ethyl acetate was added to the organic phase containing quinapril benzyl ester. The reaction mixture was stirred at 25-30⁰ C for 30 minutes and filtered to afford 40gms (70%) of the maleate salt (V); HPLC purity: >
20 97%; m.p, 139.1⁰C.

¹H NMR (CDCl₃, 200mHz) δ ppm : 7.0-7.40(m, 14.4), 6.30 (s, 2H), 5.45 (dd,2H), 5.05 (s, 2H), 4.7 (m, 2H), 4.4(m, 2H), 3.7 (m,1H), 3.05-3.45 (m,3H), 2.7-2.9 (m, 2H), 2.15-2.35 (m, 2H), 1.4-1.7(m, 3H), 1.3 (t, 3H)

25

I.R (KBr)(v, cm⁻¹): 3008, 1745, 1653, 1496, 1452, 1344, 1190, 1169, 1009, 984, 757, 652, 564.

30

Example-2

Preparation of the benzyl ester of (S, S, S)-2-[2-[(1-ethoxycarbonyl)-3-phenylpropyl)amino]-1-oxopropyl]-1,2,3,4-tetrahydro-3-isoquinoline carboxylic acid; maleic acid salt . (Quinapril benzyl ester maleic acid salt). (Method-II)

- 5 50.0gms (0.114 moles) of the para-toluene sulfonate salt of the benzyl ester of (S)-1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid (IV) was added to a mixture of dichloromethane (150ml) and water(150ml). The mixture was cooled to 5°C, and the pH of the mixture was adjusted to 8.5 by dropwise addition of 25% aqueous ammonia (10ml), at the same temperature, and stirred at 30°C for 30 minutes. The organic layer
- 10 containing the free base of (IV) was separated.
(43.4gms; 0.142moles) N[1(S)-carboethoxy-3-phenylpropyl]-(S)-alanyl carboxyanhydride (VI) was dissolved in dichloromethane(175ml) at 30°C. The organic layer containing the free base of (IV) was added in 30 minutes at 30°C. The reaction mixture was monitored was stirred for 4.0 hours for the reaction to go to completion.
- 15 Triethyl amine(5.75gms), followed by 10% sodium carbonate solution (100ml) were added to the reaction mixture at 30°C, and stirred for 4.0hours. The organic layer was separated and concentrated at 40°C. The residue was diluted with ethyl acetate (300ml) and stirred to dissolve the residue at 40°C.
- 20 Maleic acid (13.21gms) dissolved in ethyl acetate (350ml) was added at 40°C to the ethyl acetate layer containing the quinapril benzyl ester. The reaction mixture was stirred for 30 minutes at 30°C and then cooled to 0°C. The mixture was stirred for 30 minutes and filtered. The wet cake was washed with ethyl acetate and dried at 50°C/10-20mmHg for 6.0 hours, to afford 55.74 gms (76%) of the maleate salt (V); HPLC purity: > 98%; m.p,
- 25 139.1°C.

¹H NMR (CDCl₃, 200mHz) δ ppm: 7.0-7.40(m, 14.4), 6.30 (s, 2H), 5.45 (dd,2H), 5.05 (s, 2H), 4.7 (m, 2H), 4.4 (m, 2H), 3.7 (m,1H), 3.05-3.45 (m,3H), 2.7-2.9 (m, 2H), 2.15-2.35 (m, 2H), 1.4-1.7(m, 3H), 1.3 (t, 3H)

I.R (KBr)(ν , cm^{-1}): 3008, 1745, 1653, 1496, 1452, 1344, 1190, 1169, 1009, 984, 757, 652, 564.

Example -3

- 5 **Preparation of (S, S, S) 2-{2-[(1-ethoxycarbonyl-3-phenylpropyl)amino]-1-oxopropyl}-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid hydrochloride. (Quinapril hydrochloride).**

The conversion of quinapril benzyl ester maleic acid salt (V) as obtained in examples (1
10 & 2) to quinapril hydrochloride (I) is achieved by the following steps :

A. Formation of quinapril benzyl ester free base.

25gms(0.0388 moles) quinapril benzyl ester maleic acid salt (V) obtained in
Example-1 was dissolved in a mixture of 125ml water and 125ml
15 dichloromethane. The pH of the solution was adjusted between 7.5 to 8.5 by
addition of aqueous ammonia. Reaction mixture was stirred for 30 minutes, the
organic phase separated and washed with 50ml water. The organic phase was
separated and treated with activated carbon. After filtration, the organic phase was
evaporated under reduced pressure below 40°C , the free base of quinapril benzyl
20 ester was obtained.

B. Debenzylation of quinapril benzyl ester (Preparation of Quinapril hydrochloride).

The benzyl ester free base obtained in Step A is dissolved in 140ml ethanol to
which 1.0gm of 10%Pd/C and 6.0gms of 35% hydrochloric acid are added. The
25 reaction mass is subjected to catalytic hydrogenolysis with hydrogen gas at 40-60
psi pressure and $20-30^{\circ}\text{C}$. The reaction mass is filtered and the filtrate evaporated
to give crude quinapril hydrochloride. The residue is dissolved in dichloromethane
and the solvent recovered below 35°C under reduced pressure. This operation is
repeated till the water content of the residue is less than 1.0%.

Example -4**Crystallisation of quinapril hydrochloride (I) form Nitromethane**

Nitromethane (125ml) is added to the anhydrous residue of quinapril hydrochloride obtained in Step B, Example-3 and the mixture is stirred at 20-25°C for 10-15 minutes to
 5 get a clear solution. The mixture is stirred at the same temperature for 30 minutes and seeded with pure quinapril hydrochloride . The mass is cooled to 5-10°C, and stirred at the same temperature for 2 hrs. The reaction mass is then filtered at 20-25°C and the wet cake is washed with nitromethane (50ml) to give crystalline quinapril hydrochloride associated with a solvate of nitromethane, having HPLC Purity > 99% and X-ray
 10 (powder) diffraction pattern as summarized in Table-I and Fig-Ia.

IR (KBr) (ν , cm⁻¹) : 3030, 2933, 2856, 2796, 1743, 1701, 1645, 1550, 1490, 1450, 1263, 1197, 1091, 756, 729, 707.

15 Solid state ¹³C CPMAS at 10KHz: 255.04, 221.26, 212.38 207.50 175.6, 168.7, 141.9, 132.93, 127.16, 96.07, 89.25, 87.34, 62.79, 60.01, 56.71, 53.41, 47.63, 35.76, 14.58.

20	DSC (50.0-300.0°C; 05.00°C/min):	Integral :	-233.25mJ	Integral:	-245.18mJ
		Onset :	96.99°C	Onset:	151.25°C
		Peak :	107.39°C	Peak :	163.53°C
		Endset:	111.55	Endset:	172.43°C

The crystalline nitromethane solvate is then dried at 40°C / 0-5mm Hg pressure for 60 hours to give pure amorphous quinapril hydrochloride; m. p. 117-121°C; [α]_D (2% in
 25 CH₃OH): +14.92 in a yield of 65%.

¹HNMR: (CDCl₃, 200mHz) δ ppm: 10.0(bs, 1H), 8.9(bs, 1H), 7.08 (m, 9H), 5.05 (m, 1H), 4.35-4.95(m, 3H), 3.7-4.3(m, 3H), 2.9-3.4(2H), 2.45-2.85(m, 2H), 2.1-2.4(m, 2H), 1.4-1.8(dd, 3H), 1.0-1.25(m, 3H).

30 IR (KBr)(ν , cm⁻¹): 2981, 2858, 1739, 1649, 1535, 1496 1438, 1369, 1259, 1207, 750, 702.

Example -5**Crystallisation of quinapril hydrochloride from nitroethane.**

Nitroethane (125ml) is added to the anhydrous residue of impure quinapril hydrochloride obtained in step B, Example-3 and the mixture is stirred at 20-25°C for 10-15 minutes to get a clear solution. The mixture is stirred at the same temperature for 30 minutes and seeded with pure quinapril hydrochloride. The mass is cooled to 5-10°C, and stirred at the same temperature for 2 hrs. The reaction mass is then filtered at 20-25°C and the wet cake is washed with nitroethane (50ml) to give crystalline quinapril hydrochloride associated with a solvate of nitroethane, having HPLC purity >99% and X-ray (powder) diffraction pattern as summarized in Table-I and Fig-Ib.

IR (KBr) (ν , cm⁻¹): 3435, 2849, 1742, 1707, 1642, 1553, 1492, 1450, 1430, 1408, 1368, 1298, 1261, 1195, 1040, 757.

15	DSC (50.0-300.0°C; 05.00°C/min):	Integral :	-716.23mJ	Integral:	-357.15mJ
		Onset :	97.32°C	Onset:	157.48°C
		Peak :	106.64°C	Peak :	164.09°C
		Endset:	111.39	Endset:	170.80°C

The crystalline nitroethane solvate is then dried at 40°C / 0-5mm Hg pressure for 60 hours to give pure amorphous quinapril hydrochloride, m. p 117-121°C; $[\alpha]_D^{25}$ (2% in CH₃OH): +14.92 in a yield of 65%.

¹HNMR: (CDCl₃, 200MHz) δ ppm: 10.0(bs, 1H), 8.9(bs, 1H), 7.08 (m, 9H), 5.05 (m, 1H), 4.35-4.95(m, 3H), 3.7-4.3(m, 3H), 2.9-3.4(2H), 2.45-2.85(m, 2H), 2.1-2.4(m, 2H), 1.4-1.8(dd, 3H), 1.0-1.25(m, 3H).

IR (KBr)(ν , cm⁻¹): 2981, 2858, 1739, 1649, 1535, 1496, 1438, 1369, 1259, 1207, 750, 702.

Example-6**Crystallisation of quinapril hydrochloride from nitropropane**

Nitropropane (125ml) is added to the anhydrous residue of quinapril hydrochloride from step B, Example-3 and the mixture is stirred at 20-25°C for 10-15 minutes to get a clear

solution. The mixture is stirred at the same temperature for 30 minutes and seeded with pure quinapril hydrochloride. The mass is cooled to 5-10°C, and stirred at the same temperature for 2 hrs. The reaction mass is then filtered at 20-25°C and the wet cake is washed with nitropropane (50ml) to give and crystalline quinapril hydrochloride associated with a solvate of nitropropane, having HPLC purity > 99% and X-ray (powder) as summarized in Table-I and Fig-Ic.

IR (KBr) (ν , cm^{-1}): 3420, 2850, 1742, 1705, 1643, 1547, 1495, 1451, 1437, 1408, 1334, 1296, 1259, 1225, 1091, 1041, 757, 497.

10

DSC (50.0-300.0°C; 05.00°C/min):	Integral :	-325.52mJ	Integral:	-419.74mJ
	Onset :	92.34°C	Onset:	152.58°C
	Peak :	95.15°C	Peak :	159.63°C
	Endset:	97.17°C	Endset:	173.72°C

15

The crystalline material is then dried at 40°C / 0-5mm Hg pressure for 60 hours to give amorphous quinapril hydrochloride, m. p. 117-121°C; $[\alpha]_d$ (2% in CH_3OH): +14.92 in a yield of 65%.

20

^1H NMR:(CDCl_3 , 200ppm) δ ppm: 10.0(bs, 1H), 8.9(bs, 1H), 7.08 (m, 9H), 5.05 (m, 1H), 4.35-4.95(m, 3H), 3.7-4.3(m, 3H), 2.9-3.4(2H), 2.45-2.85(m, 2H), 2.1-2.4(m, 2H), 1.4-1.8(dd, 3H), 1.0-1.25(m, 3H) .

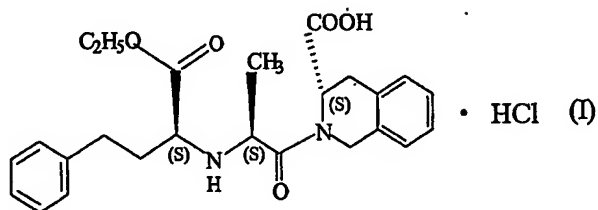
25

IR (KBr)(ν , cm^{-1}): 2981, 2858, 1739, 1649, 1535, 1496, 1438, 1369, 1259, 1207, 750, 702.

30

CLAIMS

1. A crystalline nitroalkane solvate of quinapril hydrochloride of formula (I)
 5 substantially free of impurities.



2. A crystalline nitroalkane solvate quinapril hydrochloride as claimed in claim 1
 10 wherein the nitroalkane is nitromethane, said nitromethane solvate of quinapril hydrochloride exhibiting essentially the following X-ray (powder) diffraction data.

Spacing 'd'	Relative intensity
16.2471	63.5
13.8426	55.5
11.9609	57.3
9.6467	17.2
7.9468	22.4
7.5064	31.1
7.1699	30.2
6.4095	52.4
6.0561	3.1
5.5041	20.4
5.2808	33.1
5.1767	22.8
4.8704	5.7
4.6830	34.4
4.4404	100

4.0977	30.9
3.9931	69.5
3.7747	62.8
3.5972	12.9
3.5058	22.8
3.4153	8.3
3.3558	7.0
3.2676	30.7
3.2054	7.6
3.1510	9.5
3.0854	16.5
2.9772	14.5
2.9403	17.5
2.9122	12.9
2.7798	7.6
2.6670	9.0
2.6216	6.0
2.5613	10.1
2.4650	5.2
2.3933	7.9
2.2963	4.9
2.2620	2.7
2.2290	3.0
2.1672	3.5
2.1125	3.4
2.0361	2.0
1.9911	3.3
1.9714	3.4
1.8935	2.5
1.8420	2.3
1.7917	2.0

1.7630	1.3
1.6723	1.1
1.5928	0.4
1.4683	0.5

3. A crystalline nitroalkane solvate quinapril hydrochloride as claimed in claim 1 wherein the nitroalkane is nitroethane, said nitroethane solvate of quinapril hydrochloride exhibiting essentially the following X-ray (powder) diffraction data.

5

Spacing 'd'	Relative intensity
17.4844	87.0
16.0841	67.2
12.0996	53.3
10.0860	18.8
8.1700	29.6
7.7522	41.2
7.3814	35.4
6.5032	39.9
6.0580	2.9
5.5780	35.8
5.3973	32.0
5.2776	23.7
4.9335	19.6
4.8335	32.9
4.7627	40.5
4.5635	88.9
4.5095	69.6
4.4227	36.5
4.2021	36.0
4.0818	100.0

3.8719	30.4
3.7802	48.7
3.6435	49.5
3.4889	10.4
3.3520	20.8
3.2948	36.3
3.1526	15.3
3.1132	19.1
3.0304	49.5
2.9280	19.2
2.9080	14.1
2.7933	7.9
2.6986	10.8
2.6399	9.4
2.5715	16.7
2.5194	9.6
2.4535	4.6
2.4140	12.7
2.3567	8.5
2.3093	5.5
2.2801	6.1
2.1687	6.7
2.1303	4.5
2.0332	5.0
2.0031	6.6
1.9801	3.7
1.9395	2.7
1.8918	4.2
1.8632	3.5
1.8354	3.0
1.8110	2.4

1.7812	2.1
1.7024	1.1
1.5414	0.6
1.3867	0.4

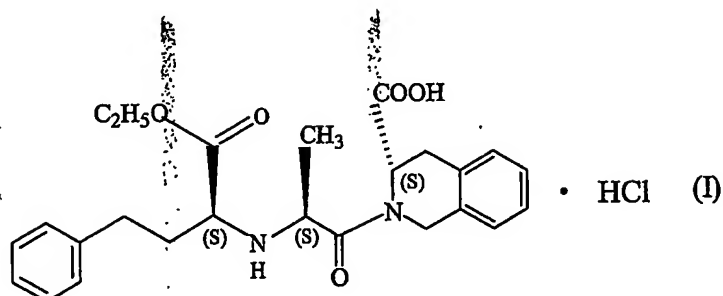
4. A crystalline nitroalkane solvate quinapril hydrochloride as claimed in claim 1 wherein the nitroalkane is nitropropane, said nitropropane solvate of quinapril hydrochloride exhibiting essentially the following X-ray (powder) diffraction data.

5

Spacing 'd'	Relative intensities
17.4327	100.0
15.4378	65.5
12.1079	49.5
10.4492	18.2
8.3626	21.3
7.9183	38.9
7.5253	27.5
6.6268	19.6
6.4583	17.2
5.6238	43.2
5.2713	27.8
5.1011	18.3
4.9816	30.7
4.8049	38.7
4.6878	52.0
4.5427	61.4
4.4480	35.3
4.3008	25.3
4.1690	68.6
4.0279	14.7

3.9483	23.9
3.7993	57.5
3.5871	15.5
3.4582	14.8
3.2960	32.0
3.1558	20.6
3.1053	26.1
3.0548	17.9
2.9559	9.2
2.9066	9.9
2.8044	10.6
2.7274	6.5
2.6357	9.4
2.5910	11.1
2.4880	5.3
2.4365	6.4
2.4053	6.0
2.3446	8.4
2.3084	5.2
2.2682	4.1
2.2075	3.0
2.1667	4.1
2.0297	13.7
2.0023	3.5
1.8765	2.2
1.8019	2.6
1.4337	2.2
1.4005	0.4

5. A process for preparation of quinapril hydrochloride of formula (I) substantially free of impurities

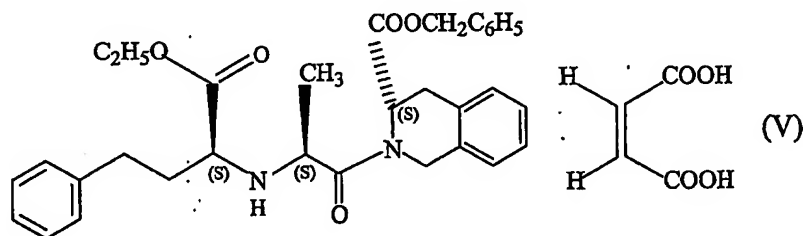


5

which comprises the steps of :

- a) adjusting the pH between 7.5-8.5 of a solution of the benzyl ester maleate salt of quinapril of formula (V) in a mixture of water and an organic solvent to obtain free base compound of formula (V),

10



- b) subjecting the free base of compound (V) thus obtained in step a) to catalytic hydrogenation in an alcoholic solvent in the presence of concentrated hydrochloric acid or hydrogen chloride dissolved in an alcoholic solvent and in the presence of catalytic amounts of Pd/C to obtain a residue containing formula (I),

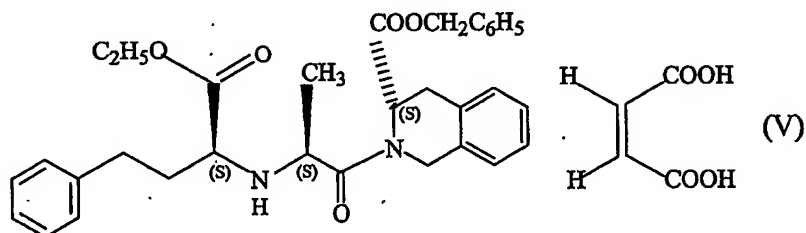
15

- c) crystallization of the residue containing compound of formula (I) thus obtained by evaporating the alcoholic solvent from step b) from a nitroalkane solvent to give crystalline quinapril hydrochloride, associated with a solvate of the nitroalkane solvent, and

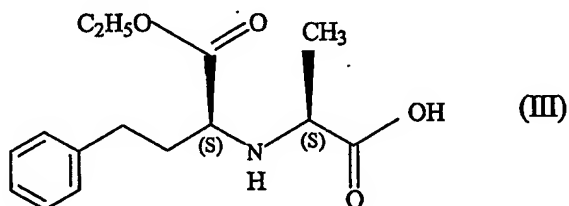
20

d) drying the crystalline quinapril hydrochloride nitroalkane solvate obtained in step c) at a temperature between 40° C and 45° C under vacuum to give pure quinapril hydrochloride of formula (I).

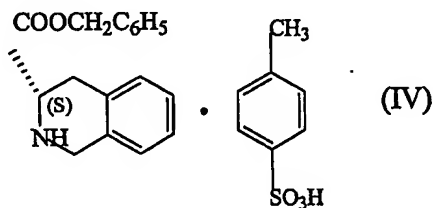
5 6. A process according to Claim 5, wherein the benzyl ester maleate salt of quinapril of formula (V),



is obtained by reacting the acid halide of compound of formula (III),



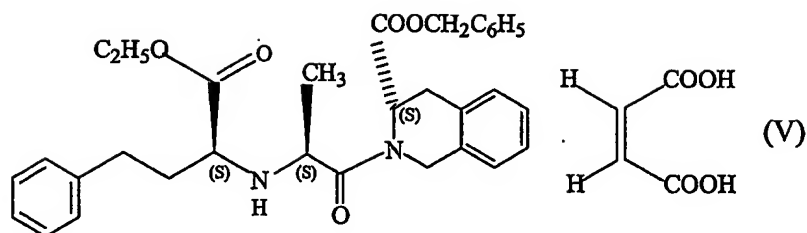
with compound of formula (IV)



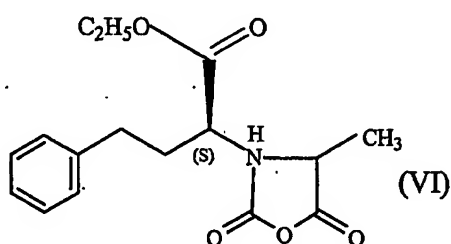
in an organic solvent and in the presence of a base followed by addition of maleic acid.

7. A process according to Claim 5, wherein the benzyl ester maleate salt of quinapril of formula (V),

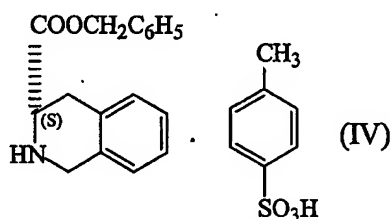
37



is obtained by reacting the compound of formula (VI),



5 with a compound of formula (IV),



in an organic solvent and in the presence of a base followed by addition of maleic acid.

10 8. A process according to step (a) of Claim 5, wherein the organic solvent is selected from a chlorinated hydrocarbon.

9. A process according to Claim 8, wherein the chlorinated hydrocarbon is selected from dichloromethane, 1,2-dichloroethane and chloroform.

15

10. A process according to step (a) of Claim 5, wherein the pH is adjusted using aqueous ammonia.

11. A process according to step (b) of Claim 5, wherein the alcoholic solvent is selected from methanol, ethanol, n-propanol and isopropanol.
12. A process according to step (c) of Claim 5, wherein the nitroalkane solvent is selected
5 from nitromethane, nitroethane and nitropropane.
13. A process according to step (c) of Claim 5, wherein the compound obtained in Step (b) is dissolved in the nitroalkane solvent at room temperature.
- 10 14. A process according to step (c) of Claim 5, wherein the crystallization is carried out a temperature ranging from -15°C to $+15^{\circ}\text{C}$.
- 15 15. A process according to Claim 14, wherein the crystallization is carried out a temperature ranging from 0°C to $+10^{\circ}\text{C}$.
16. A process according to step (d) of Claim 5, wherein the drying is carried out under a vacuum of 5-10 mm Hg and lasts from about 60 to about 70 hours.
17. A process according to claim 6, wherein the acid halide is acid chloride or acid
20 bromide.
18. A process according to Claim 6, wherein the organic solvent is selected from a chlorinated hydrocarbon or an alkyl acetate.
- 25 19. A process according to Claim 18, wherein the chlorinated hydrocarbon is selected from dichloromethane, 1,2-dichloroethane and chloroform.
20. A process according to Claim 18, wherein the alkyl acetate is selected from methyl acetate, ethyl acetate and butyl acetate.
- 30 21. A process according to Claim 6, wherein the base is an organic base.

22. A process according to Claim 21, wherein the organic base is selected from triethylamine, diethylamine, tertiary butylamine, and dicyclohexylamine.
23. A process according to Claim 7, wherein the organic solvent is selected from a
5 chlorinated hydrocarbon or an alkyl acetate.
24. A process according to Claim 23, wherein the chlorinated hydrocarbon is selected from dichloromethane, 1,2-dichloroethane and chloroform.
- 10 25. A process according to Claim 23, wherein the alkyl acetate is selected from methyl acetate, ethyl acetate and butyl acetate.
26. A process according to Claim 7, wherein the base is an organic base.
- 15 27. A process according to Claim 7, wherein the base is an inorganic base.
28. A process according to Claim 26, wherein the organic base is selected from triethylamine, diethylamine, tertiary butylamine, and dicyclohexylamine.
- 20 29. A process according to Claim 27, wherein the inorganic base is selected from sodium carbonate, sodium bicarbonate, potassium carbonate, potassium bicarbonate.

Fig-1a: X-Ray (powder) diffraction pattern of quinapril hydrochloride crystallized from nitromethane.

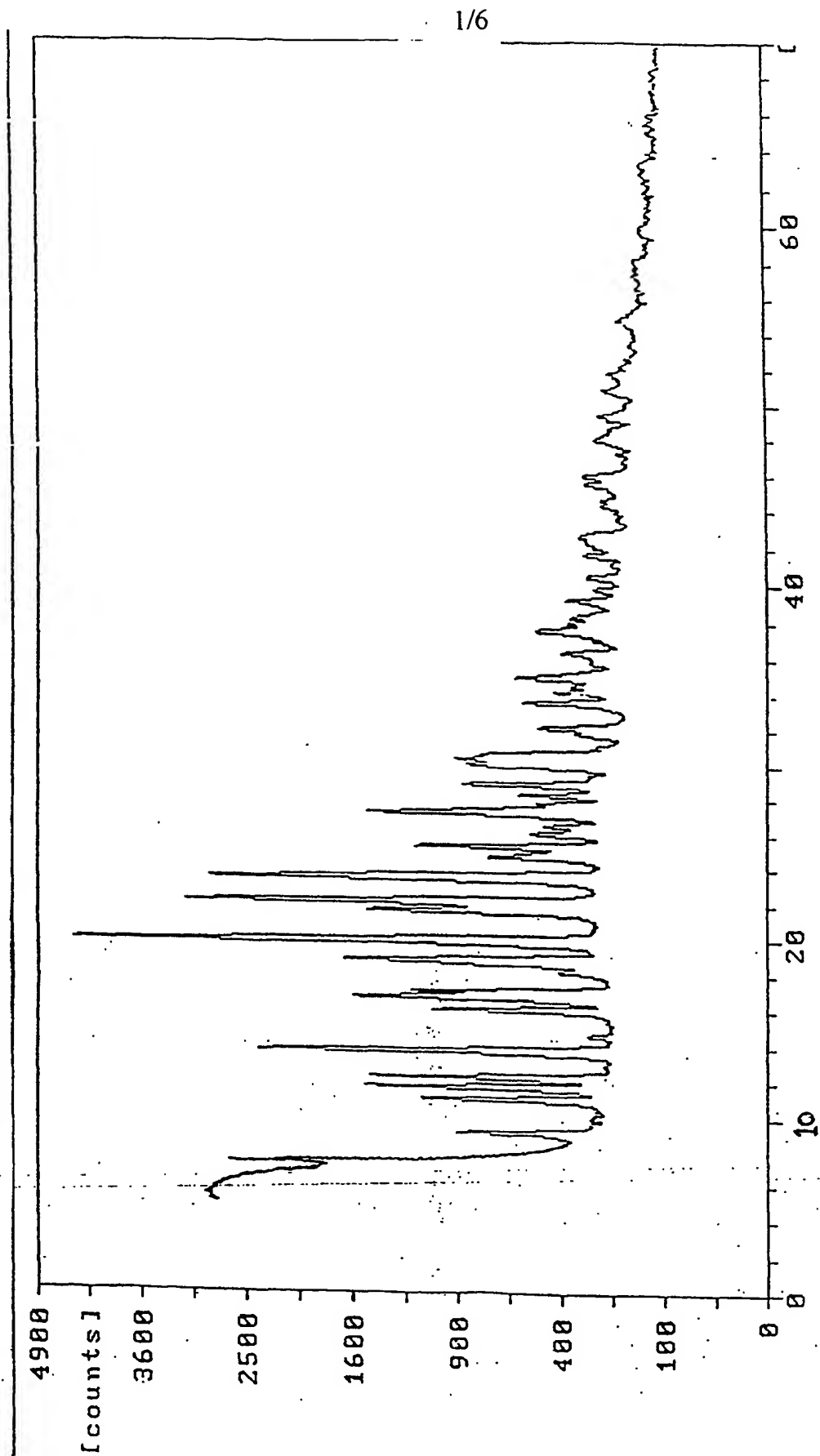


Fig-1b: X-Ray (powder) diffraction pattern of quinapril hydrochloride crystallized from nitroethane.

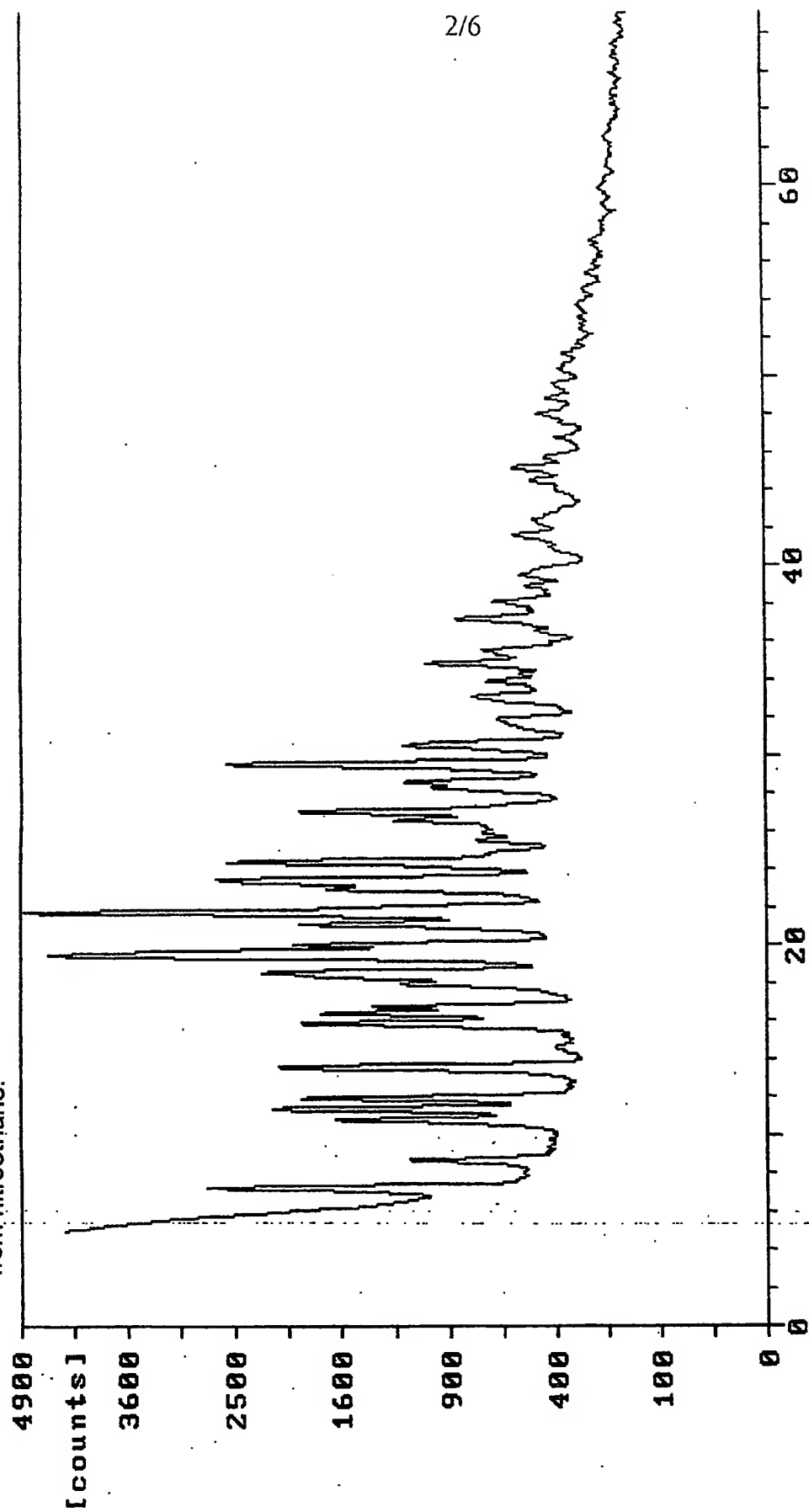


Fig-1c: X-Ray (powder) diffraction pattern of quinapril hydrochloride crystallized from nitropropane.

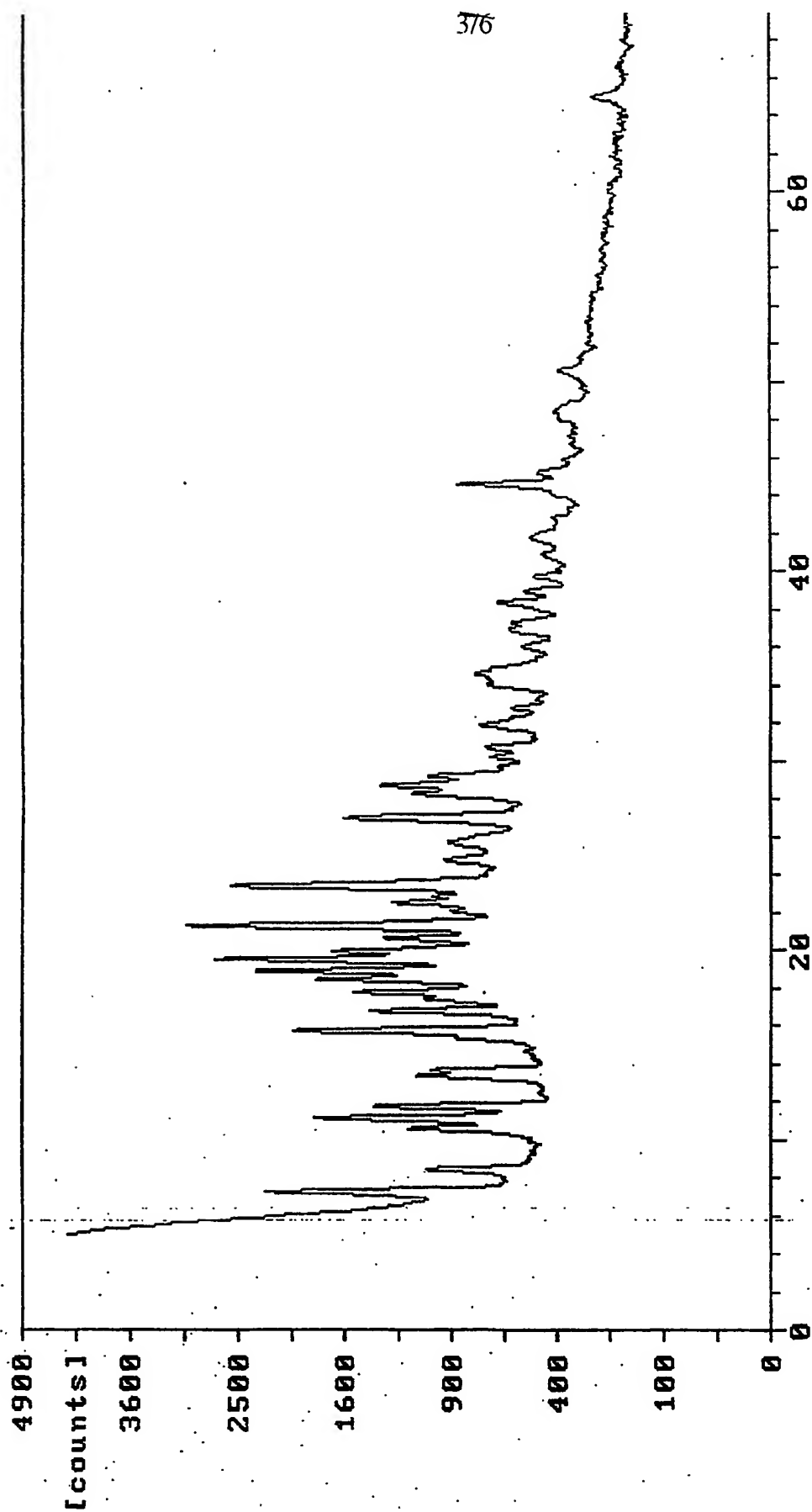
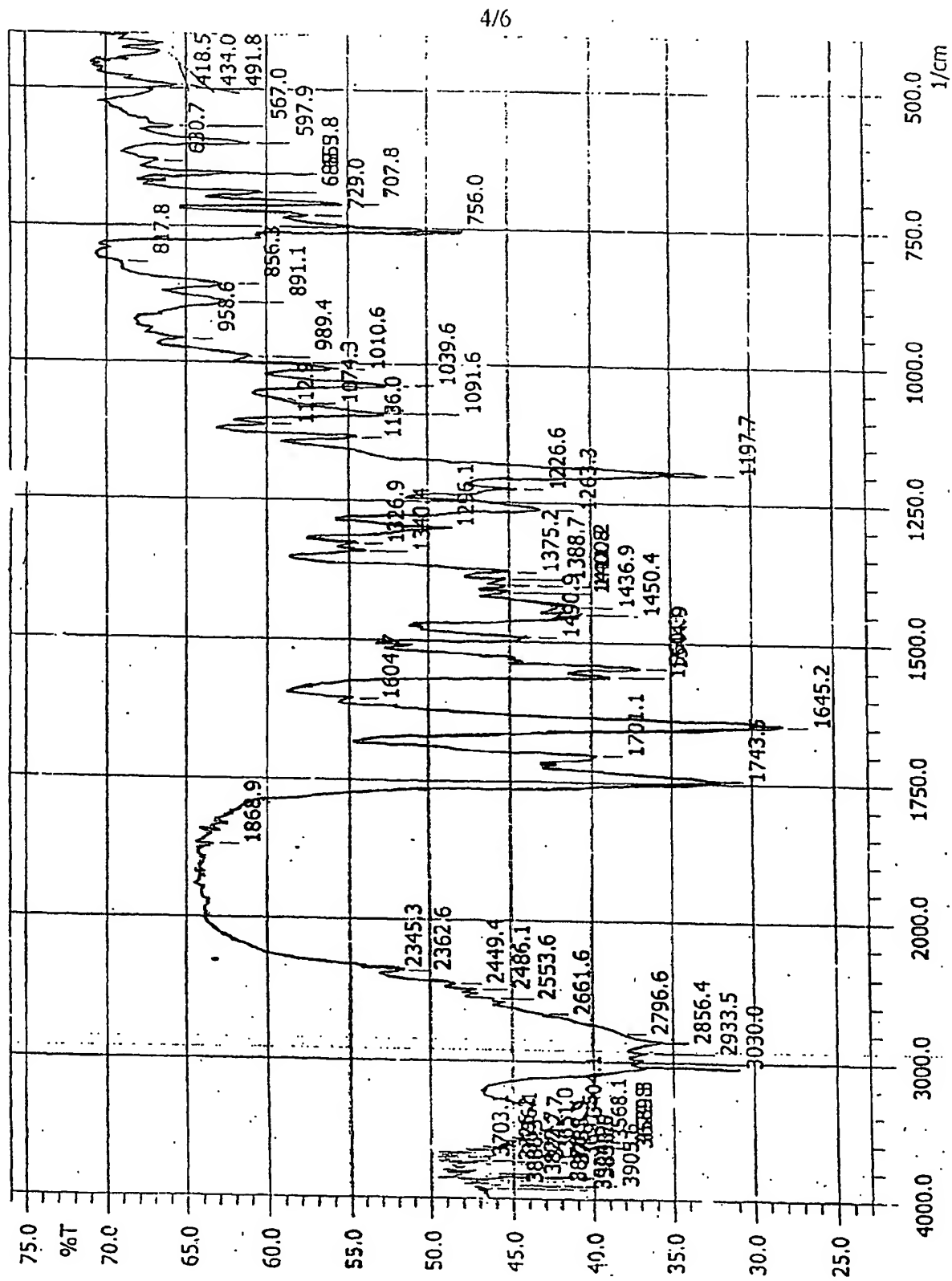
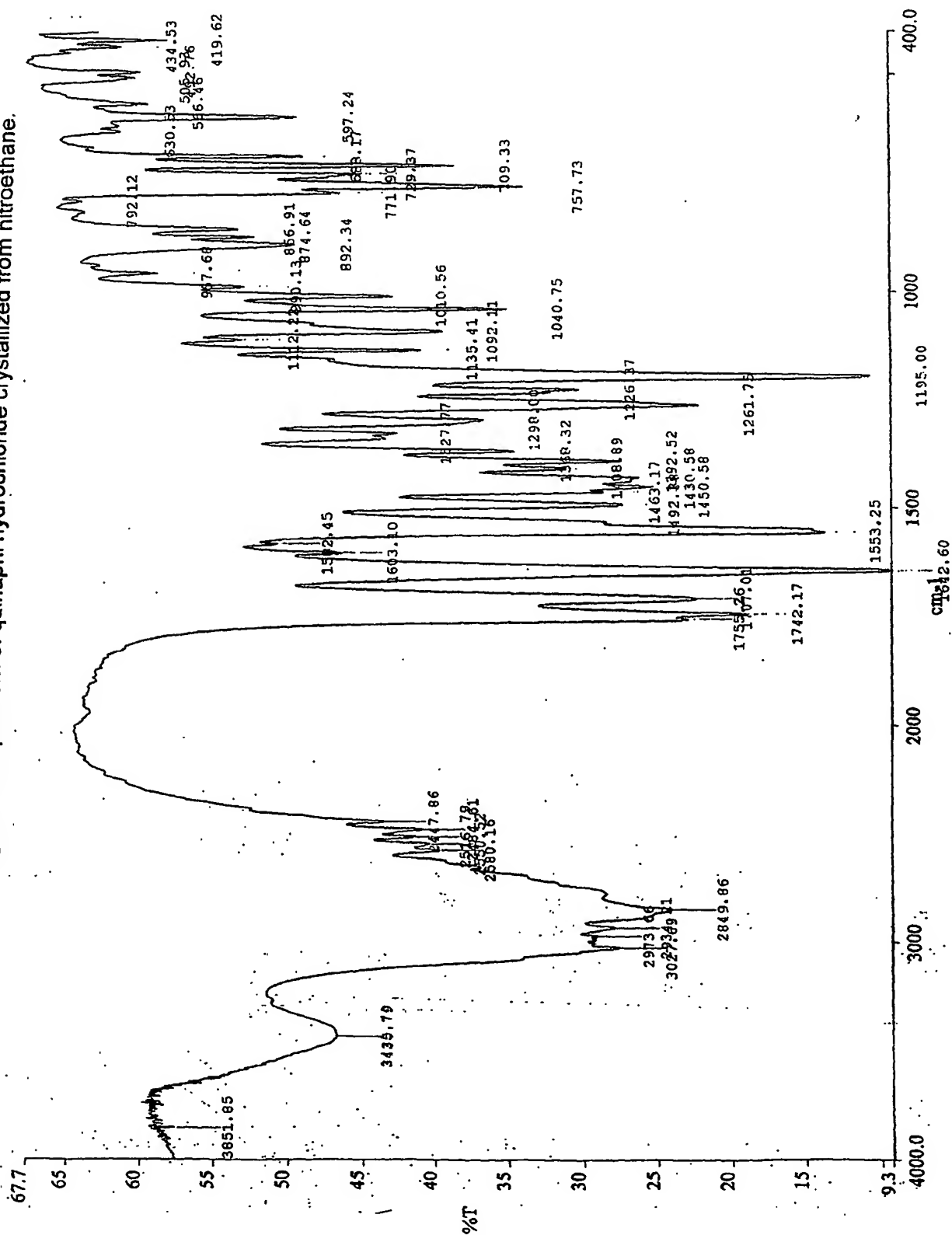


Fig-11a: I.R. spectrum of quinapril hydrochloride crystallized from nitromethane.



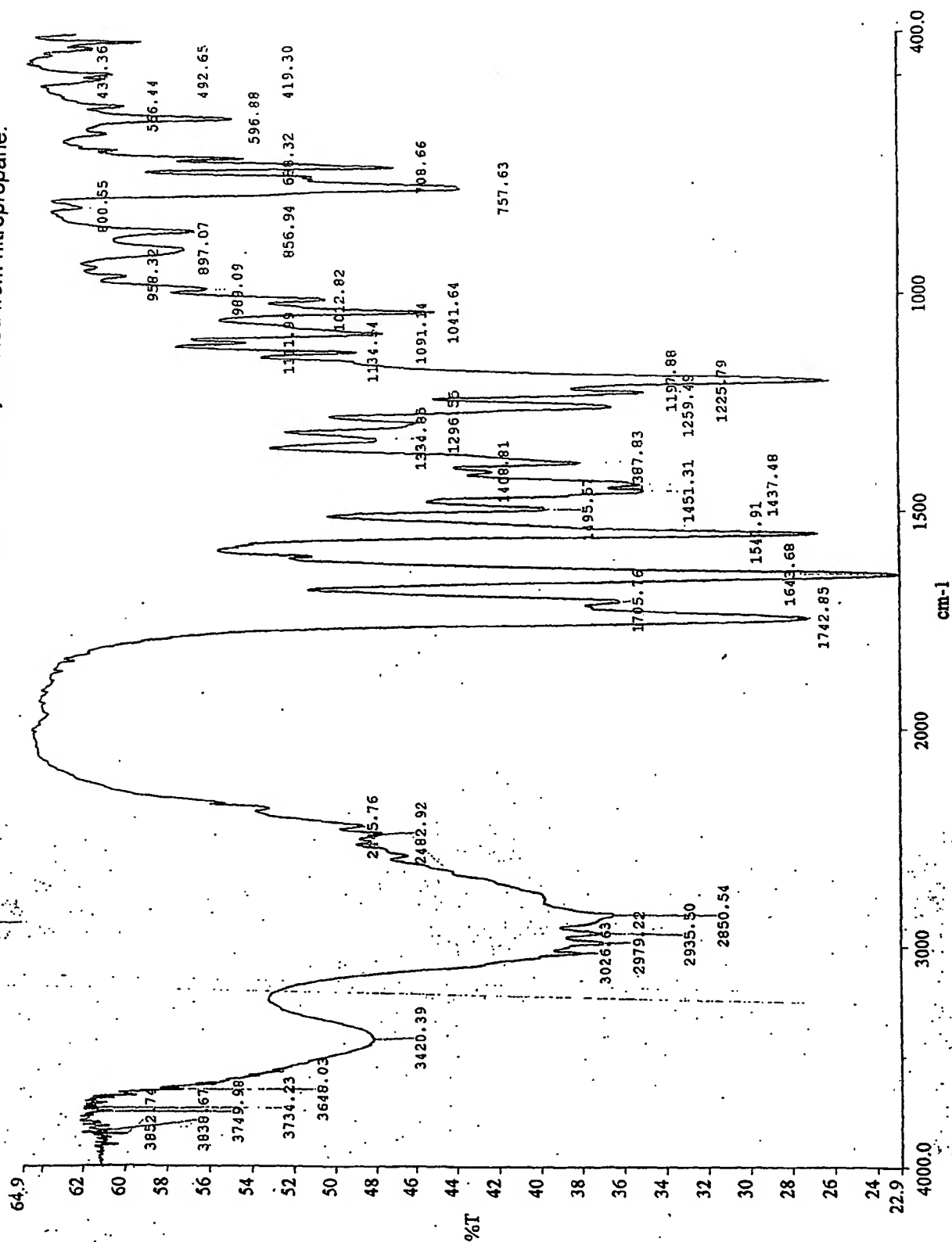
5/6

Fig-11b: I.R. spectrum of quinapril hydrochloride crystallized from nitroethane.



6/6

Fig-11c: I.R. spectrum of quinapril hydrochloride crystallized from nitropropane.



INTERNATIONAL SEARCH REPORT

Internal application No
PCT/IN 02/00235

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C07D217/26

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the International search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, BEILSTEIN Data, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	EP 0 285 992 A (WARNER LAMBERT CO) 12 October 1988 (1988-10-12) cited in the application the whole document	1-29
Y	EP 0 992 495 A (ESTEVE QUIMICA SA) 12 April 2000 (2000-04-12) cited in the application the whole document	1-29
A	US 4 344 949 A (HOEFLE MILTON L ET AL) 17 August 1982 (1982-08-17) cited in the application the whole document	1-29

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex. ---

* Special categories of cited documents:

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

30 June 2003

Date of mailing of the international search report

04/07/2003

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Scruton-Evans, I

INTERNATIONAL SEARCH REPORT

International Publication No

PCT/IN 02/00235

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
EP 0285992	A	12-10-1988	US 4761479 A	02-08-1988
			AT 62229 T	15-04-1991
			AU 605555 B2	17-01-1991
			AU 1238388 A	29-09-1988
			CA 1291999 C	12-11-1991
			DE 3862227 D1	08-05-1991
			EP 0285992 A1	12-10-1988
			GR 3001736 T3	23-11-1992
			JP 2659990 B2	30-09-1997
			JP 63258459 A	25-10-1988
			KR 9607087 B1	27-05-1996
			NZ 223671 A	26-10-1990
			PH 24986 A	26-12-1990
			ZA 8801426 A	25-10-1989
EP 0992495	A	12-04-2000	ES 2122941 A1	16-12-1998
			AU 730140 B2	01-03-2001
			AU 7336598 A	30-12-1998
			EP 0992495 A1	12-04-2000
			JP 2002502386 T	22-01-2002
			WO 9854149 A1	03-12-1998
			ZA 9804466 A	01-12-1998
US 4344949	A	17-08-1982	AT 25974 T	15-04-1987
			AT 26120 T	15-04-1987
			AU 563683 B2	16-07-1987
			AU 5299186 A	17-07-1986
			AU 551239 B2	24-04-1986
			AU 7541681 A	06-05-1982
			CA 1341330 C	01-01-2002
			CA 1331614 C	23-08-1994
			CA 1331615 C	23-08-1994
			DD 201787 A5	10-08-1983
			DE 3176010 D1	23-04-1987
			DE 3176029 D1	30-04-1987
			DK 436081 A ,B,	04-04-1982
			EP 0049605 A1	14-04-1982
			EP 0096157 A2	21-12-1983
			ES 8301925 A1	01-04-1983
			FI 813033 A ,B,	04-04-1982
			FI 881985 A ,B,	27-04-1988
			GR 75353 A1	13-07-1984
			HK 43389 A	07-06-1989
			IE 52666 B1	20-01-1988
			IE 52665 B1	20-01-1988
			IL 63806 A	28-02-1985
			JP 1653003 C	30-03-1992
			JP 3016359 B	05-03-1991
			JP 57088164 A	01-06-1982
			KR 8801326 B1	25-07-1988
			LU 88321 A9	04-05-1994
			MX 155144 A	01-02-1988
			NO 813359 A ,B,	05-04-1982
			PH 17740 A	23-11-1984
			SG 15689 G	09-06-1989
			US 4532342 A	30-07-1985
			HU 183602 B	28-05-1984
			NZ 198542 A	20-03-1985

INTERNATIONAL SEARCH REPORT

International Application No
PCT/IN 02/00235

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4344949	A	SU 1148560 A3 ZA 8106332 A	30-03-1985 29-09-1982
<hr/>			